

Client: ██████████
Accession: ██████████ **Spec Type:** SWAB
Initial Report Date: 12/19/2020
Final Report Date: 12/19/2020

1 Industry Dr., Henderson, NC 27537
Phone: 919-351-6256, Fax: 252-572-4595
CLIA ID: 34D2141858



Order Information

Patient Information:
OPPEDAHL, CARL
Date of Birth: ██████████
Gender: Male Race: W
Patient ID: ██████████
Phone: 970-██████-6088
Collected: 12/17/2020 11:08
Address: ██████████

Account Information:
Acct #: ██████████
Clinician: ██████████
Location: ██████████
Collector: ██████████
Requisition: ██████████
Received: 12/18/2020 04:38

Test Information:
720100: SARS-CoV-2

Diagnosis Codes: Diagnosis Codes Not Provided

Medications: No Medications Indicated

Comments:

	RESULT	(Abnormal)	FLAG	UOM	REFERENCE RANGE	LOC
SARS-CoV-2	Not Detected					

Negative (Not Detected)

The SARS-CoV-2 test is intended for the presumptive qualitative detection of nucleic acid from SARS-CoV-2 in upper respiratory specimens. A Negative result does not exclude infection caused by SARS-CoV-2 and the results must be correlated with clinical presentation and evaluated in the context of other laboratory and epidemiologic data.

The COVID-19 RT-qPCR test is a Laboratory Developed test (LDT) not approved or cleared by the FDA and has been approved for Emergency Use Authorization (EUA). This test has been authorized by FDA under an Emergency Use Authorization (EUA). This test is only authorized for the duration of time the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection of SARS-CoV-2 virus and/or diagnosis of COVID-19 infection under section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.

(1) Test Performed at: Mako Medical - Henderson
1 Industry Dr. Henderson, NC, 27537
Director: Chad R. Rund, D.O., FCAP 34D2141858

*** Final Report ***

Member: Carl Oppedahl
Date of birth: [REDACTED]
Sex: M
Primary care physician: [REDACTED]
Date printed: 12/18/2020

For general information about a test procedure, click the "About this test" link above.

To see more information about a test result, select the "Details" tab. To compare test results over time, click "Past results" or "Graph of past results."

E-mail your doctor or schedule a video or phone visit if you would like to discuss these results.

Result Information

Status: Final result
(Collected: 12/18/2020
1:44 PM)

Provider Status: Ordered

12/18/2020 7:33 PM

Component

SARS-COV-2 (COVID-19) IgG, IgM Antibody Interpretation

Reactive

Narrative

A REACTIVE result means your test DID detect COVID-19 IgG, IgM antibodies